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Immune response of calves inoculated with proteins of *Anaplasma marginale* bound to an immunostimulant complex

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**Abstract**

Despite our current knowledge of the immunology, pathology, and genetics of *Anaplasma marginale*, prevention in cattle is currently based on old standbys, including live attenuated vaccines, antibiotic treatment, and maintaining enzootic stability in cattle herds. In the present study, we evaluated the use of an immunostimulant complex (ISCOMATRIX) adjuvant, associated with a pool of recombinant major surface proteins (rMSP1a, rMSP1b, rMSP4 and rMSP5) to improve the humoral immune response triggered in calves mainly by IgG2. Ten calves were divided in three groups: 4 calves were inoculated with the ISCOMATRIX/rMSPs (G1); 2 calves were inoculated with ISCOMATRIX adjuvant (G2); and 4 calves received saline (G3). Three inoculations were administered at 21-day intervals. In G1, the calves showed significant increases in total IgG, IgG1 and IgG2 levels 21 days after the second inoculation, compared to the control group (p < 0.05), and G1 calves remained above the cut-off value 28 days after the third inoculation (p < 0.05). The post-immunized sera from calves in G1 reacted specifically for each of the rMSPs used. In conclusion, the ISCOMATRIX/rMSPs induced antigen-specific seroconversion in calves. Therefore, additional testing to explore the protection induced by rMSPs, both alone and in conjunction with proteins previously identified as subdominant epitopes, is warranted.

Keywords: Bovine anaplasmosis, vaccine, MSP1a, MSP1b, MSP4, MSP5.

**Resumo**

Apesar dos avanços da imunologia, patologia e genética de *Anaplasma marginale*, a prevenção em bovinos ainda é baseada nas vacinas vivas atenuadas, na terapia com antibiótico e estabilidade enzoótica dos rebanhos bovinos. No presente estudo, avaliou-se o uso de um complexo imunoestimulante (ISCOMATRIX), associado às proteínas recombinantes de superfície (rMSP1a, rMSP1b, rMSP4 e rMSP5) para melhorar a resposta imune humoral desencadeada em bezerros, principalmente por IgG2. Dez animais foram divididos em três grupos: 4 bezerros foram inoculados com o ISCOMATRIX/rMSPs (G1); 2 bezerros foram inoculados com ISCOMATRIX adjuvante (G2) e 4 bezerros receberam salina (G3). Três doses vacinais foram administradas em intervalos de 21 dias. No G1, os bezerros apresentaram aumentos significativos nos níveis de IgG total, IgG1 e IgG2 21 dias após a segunda inoculação, em comparação com o grupo de controle (p < 0.05). No G1, os bezerros apresentaram aumentos significativos nos níveis de IgG total, IgG1 e IgG2 21 dias após a segunda inoculação, em comparação com o grupo de controle (p < 0.05). Os soros pós-imunização de bezerros do G1 reagiram especificamente com cada uma das rMSPs utilizadas. Em conclusão, o ISCOMATRIX/rMSPs induziu soroconversão anti-gênero-específica em bezerros. Portanto, se justifica a realização de ensaios adicionais para explorar a proteção induzida pela rMSPs, tanto sozinhas como em conjunto com novas proteínas identificadas com epitopos subdominantes.

Palavras-chave: Anaplasmos bovina, vacina, MSP1a, MSP1b, MSP4, MSP5.

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Introduction

*Anaplasma marginale*, of the family Anaplasmataceae, is an obligate intraerythrocytic bacterium that causes bovine anaplasmosis (DUMLER et al., 2001). Transmission of *A. marginale* to cattle occurs biologically by ticks and mechanically by biting flies and by blood-contaminated fomites. Both male ticks and cattle hosts can become persistently infected with *A. marginale*, serving as reservoirs of infection (KOCAN et al., 2010). This disease is responsible for serious economic losses, mainly in tropical and subtropical regions, due to fever and hemolysis with associated severe anemia, which results in high rates of morbidity and mortality in susceptible animals (RICHEY; PALMER, 1990). The use of vaccines has been suggested as an alternative to control the disease; however, the commercially available vaccines consist of live or dead organisms, and the present limitations consist of the need for cryopreservation, insufficient efficacy, and the possibility of induction of iso-antibodies against erythrocytes that are transferred by colostrum, resulting in isooerythrolysis at birth (KUTTLER, 1984).

Six major surface proteins (MSPs) of *A. marginale* have been well characterized as MSP1a, MSP1b, MSP2, MSP3, MSP4, and MSP5 (PALMER et al., 1989; TEBELE et al., 1991; OBERLE et al., 1993; ALLEMAN; BARBET, 1996; VISSER et al., 1992). These proteins are responsible for the interaction of *A. marginale* with host cells, and they include adhesion proteins and MSPs from multigene families (PALMER et al., 1999, KOCAN et al., 2010). MSP1a, MSP4, and MSP5 are present and have been conserved in many Brazilian isolates of *A. marginale* (KANO et al., 2002), suggesting their potential use as components for a recombinant protein vaccine for anaplasmosis.

Recently, in addition to the well-characterized major surface proteins MSP1 through MSP5, other proteins, classified as immunologically ‘subdominant’ antigens, have been identified as protective antigens of *A. marginale* (KANO et al., 2002), suggesting their potential use as components for a recombinant protein vaccine for anaplasmosis.

Expression of MSP genes and purification of recombinant MSPs proteins

*Escherichia coli* BL21 Star (DE3) One Shot cells (Invitrogen Life Technologies, Sao Paulo, BRA) were transformed with the recombinant plasmids pET102-MSP1a, pET102-MSP1b, pET102-MSP4 and pET102-MSP5 to produce recombinant MSP1a and MSP1b (TAMEKUNI et al., 2009), MSP4 (KAWASAKI et al., 2007a) and MSP5 (MARANA et al., 2009) as a fusion product with 6His. The transformants were grown in Luria Bertani (LB) medium to an optical density (OD) of OD$_{590nm}$ 0.7, and Isopropylthio-D-Galactoside (IPTG) was added to a final concentration of 1 mM. Bacteria were harvested at 5 h post-induction by centrifugation; pellets were re-suspended in lysis buffer, and expression was analyzed from soluble and insoluble fractions on 12% SDS-PAGE gels. The culture of induced bacteria was centrifuged and lysed by sonication. The suspension was then centrifuged and the supernatant utilized for protein purification using the Ni-NTA resin columns method (Qiagen, Valencia, CA, USA); protein concentrations were determined as previously described (BRADFORD, 1976). The induced rMSP1a and rMSP1b proteins (74 kDa and 100 kDa) were purified under native conditions (TAMEKUNI et al., 2009), and MSP4 and MSP5 (47 kDa and 31 kDa) were purified as described by Kawasaki et al. (2007a).

**ISCOMATRIX/MSP preparation**

The ISCOMATRIX adjuvant was prepared and examined, using negative staining transmission electron microscopy, to confirm the formation of characteristic vesicles, as previously described (KAWASAKI et al., 2007b). ISCOMATRIX/rMSPs were prepared by a previously described technique (KAWASAKI et al., 2007b). The ISCOMATRIX/rMSPs were produced by the addition of 200 μg of each rMSP (rMSP1a, rMSP1b, rMSP4 and rMSP5) (1:1) to the adjuvant.

**Inoculation**

The animals were kept in three separated cattle pens and were fed silage and concentrate twice per day and water *ad libidum*. The calves were sprayed weekly with deltamethrin (Butox, Chemo)
to keep them free from ticks and biting flies. At the start of the experiment, all of the calves were approximately 240 days old and were determined to be free of *A. marginale* infection by PCR (LEW et al., 2002).

Ten calves were randomly divided into three groups: Four calves in group G1 were inoculated with the ISCOMATRIX/rMSPs, two calves in group G2 were inoculated with the ISCOMATRIX adjuvant, and four calves in group G3 were inoculated with saline. The inoculations were administered intramuscularly at days 0, 21, and 42.

**Monitoring of the experimental groups**

Before each inoculation, the animals were bled, the serum was separated, the rectal temperature was recorded, the packed cell volume (PCV) was determined, and the titer of antibodies was analyzed by indirect enzyme immunoassay (iELISA).

**Antibody assays**

Detection of anti-*A. marginale* total IgG, IgG1 and IgG2 antibodies was performed by iELISA. Optimal dilutions were established using checkerboard titrations, with dilutions of sera, antigen, and conjugates. Into each well of ELISA plates (Nunc-Immuno™ Maxisorp, Nunc, Roskilde, Denmark) were added 100 µL of *A. marginale* (strain PR1) initial bodies (10 µg/mL) diluted in sodium carbonate-bicarbonate buffer (0.05 M, pH 9.6). After overnight incubation at 4 °C, the plates were washed with PBS containing 0.05% Tween-20 (GE Healthcare, USA) (PBS-Tween pH 7.4). For total IgG, the plates were blocked for 1 hour at 37 °C using 200 µL of 8% non-fat dry milk and were washed five times with PBS-Tween (pH 7.4) in an automatic microplate washer (Bio-Rad ImmunoWash, California, USA). For IgG1 and IgG2 antibodies, the plates were blocked using 200 µL of 8% horse serum. For total IgG, serum samples were diluted (1:400) in PBS-Tween (pH 7.4) plus 5% rabbit normal sera, and 100 µL were added in duplicate to each well and then incubated at 37 °C for 90 minutes. For IgG1 and IgG2 antibodies, serum samples were diluted (1:200) in PBS-Tween (pH 7.4) plus 5% horse serum, and 100 µL was added in duplicate to each well and incubated at 37 °C for 1 hour. Positive and negative control sera were included in each plate. ELISA plates were washed five times with PBS-Tween (pH 7.4) in an automatic microplate washer (Bio-Rad ImmunoWash). Rabbit anti-bovine IgG alkaline phosphatase conjugate (Sigma Aldrich Inc., St. Louis, MO, USA), diluted 1:20,000 in PBS-Tween (pH 7.4), was added (100 µL) to each well and incubated at 37 °C for 90 minutes. For IgG1 and IgG2 antibodies, two secondary antibodies were used: horseradish peroxidase-labeled sheep anti-bovine IgG1 (Bethyl™ Laboratories, Montgomery, TX, USA) and sheep anti-bovine IgG2 (Bethyl™ Laboratories), diluted 1:10,000 and 1:15,000, respectively, in PBS-Tween (pH 7.4). Then, they were added (100 µL) to each well and incubated at 37 °C for 1 hour. The plates were then washed five times with PBS-Tween (pH 7.4) in an automatic microplate washer (Bio-Rad ImmunoWash), and 100 µL of o-phenylenediamine (OPD) (Sigma-Aldrich) solution was added in a concentration of 0.4 mg/mL using appropriate diluents with hydrogen peroxide. The reaction was interrupted by adding 50 µL of 1N HCl, and the OD reading at 490 nm was obtained using an ELISA reader (iMark™ Microplate Absorbance Reader, Bio-Rad Inc., Hercules, CA, USA).

The absorbance values were estimated, and the OD values were calculated as previously described (GARCIA et al., 2006). A serum was considered to be positive when OD sample > OD mean from negative control sera (n=10) ± 3 SDs (standard deviation) from the negative control.

**Western blot analysis**

The recombinant proteins rMSP1a, rMSP1b, rMSP4 and rMSP5 were electrophoresed and transferred, and the membranes were blocked, as previously described (KAWASAKI et al., 2007b). The membranes were incubated for 1 hour with a post-inoculation pool of serum from G1, diluted (1:500) in PBS-Tween plus 5% non-fat dry milk. The membranes were washed and incubated with peroxidase-labeled protein G (1:1,000) for 1 hour at room temperature. The peroxidase activity was demonstrated using 3,3’-diaminobenzidine (DAB) (ACROS-Organics, New Jersey, USA). Protein molecular weight markers (BenchMark™ Invitrogen Life Technologies, Carlsbad, CA, USA) were used as standards.

**Statistical analysis**

The data were first tested for normality and homogeneity of variances, and if they did not present normal distribution, they were analyzed by nonparametric statistical tests. Differences between groups at each moment were verified by the Kruskal-Wallis test, followed by Dunn’s multiple comparison test. Statistical analysis was considered significant when p < 0.05. Data were analyzed using BioEstat software, version 5.0 (AYRES et al., 2007).

**Results**

**Seroconversion**

The data for total IgG obtained by iELISA are shown in Figure 1. Calves from G1 showed a significant increase in total IgG levels 21 days after the second inoculation, compared to the control group (p = 0.0368), and the levels remained above the cut-off value 28 days after the third inoculation (p = 0.0379). Animals from G2 and G3 presented median OD values for total IgG below the cut-off value until 28 days after the third inoculation (Figure 1).

The data for IgG1 obtained by iELISA are shown in Figure 2. Calves from G1 showed a significant increase in IgG1 levels 21 days after the second inoculation, compared to the control group (p = 0.0379), and the levels remained above the cut-off value 28 days after the third inoculation (p = 0.0364). Animals from G2 and G3 presented median OD values for IgG1 below the cut-off value until 28 days after the third inoculation (Figure 2).
The data for IgG2, obtained by indirect ELISA, are shown in Figure 2. Animals from G1 presented humoral responses 21 days after the second inoculation, compared to the control group \((p = 0.0322)\), and remained above the cut-off value 28 days after the third inoculation \((p = 0.0322)\). Calves from G2 and G3 presented median OD values for IgG2 below the cut-off value during the entire study (Figure 2).

**Western blot analysis**

Western blot results, using a pool of sera of calves from G1, are shown in Figure 3. The post-inoculation sera of animals from G1 reacted specifically for each rMSP used: rMSP1a (74 kDa), rMSP1b (100 kDa), rMSP4 (47 kDa) and rMSP5 (31 kDa) (Figure 3). None of the pre-inoculation and post-inoculation sera of calves from G2 and G3 exhibit any reactivity.

**Discussion**

Prevention of anaplasmosis in cattle is currently limited and is based on the use of several classic methods: live vaccines, which are not available worldwide; antibiotic therapy, which is expensive; and acaricide control of the tick vector, which can induce resistance (SUAREZ; NOH, 2011). Protection against bovine anaplasmosis has been the target of several studies (LOPEZ et al., 2007; PALMER et al., 2011; DARK et al., 2011; LASMAR et al., 2012). Although previous studies have shown that the \(A. \) marginale subsp. centrale provides some protection against \(A. \) marginale strains (ANZIANI et al., 1987, VIDOTTO et al., 1998), others have shown partial protection with low efficacy against strains from Australia, South America, and Africa (BRIZUELA et al., 1998; TURTON et al., 1998, BOCK; DE VOS, 2001). In addition, subsequent reinfection with a high parasite load can not prevented (SHKAP et al., 2008). The basis of immunization against \(A. \) marginale with the \(A. \) marginale subsp. centrale is induction of an IgG2 response against a set of common outer membrane proteins (OMPs) expressed by both parasites (AGNES et al.,
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2011). However, antibodies are not induced to MSPs 1-3, as there is low sequence conservation, nor to MSPs 4 or 5, despite high sequence identity.

These observations and the observation that the use of subunit vaccines indicates an induction of partial immune responsiveness to challenge (PALMER; McELWAIN, 1995), inspired the present study. Here, ISCOMATRIX was used as an adjuvant, together with rMSPs (rMSP1a, rMSP1b, rMSP4, and rMSP5) of *A. marginale*. ISCOMATRIX is a cage-like structure composed by saponin (purified fraction of *Quillaja* saponin), cholesterol, and phospholipids, and it possesses immunomodulatory and antigen delivery capabilities and facilitates antigen presentation to antigen-presenting cells, such as DCs, induction of DC maturation, recruitment of immune cells to draining lymph nodes via cytokine and chemokine induction, and activation of both the innate and adaptive immune systems (MORELLI et al., 2012). Heifers vaccinated with *Staphylococcus aureus* plus ISCOMATRIX showed significantly higher levels of anti-bacterin, IgG and IgG2 in sera than animals immunized with Al(OH)3 (CAMUSSONE et al., 2013).

In the present study, calves immunized with ISCOMATRIX/rMSPs showed a peak of antibodies 21 days after the second inoculation, with antibody levels for total IgG, IgG1 and IgG2 remaining significantly above the cut-off value until 28 days after the third inoculation. However, a decrease in antibody levels for IgG3 subtypes was observed 28 days after the third inoculation, suggesting that two inoculations might be adequate to raise antibody titers. Significant increases were also observed in IgG1 and IgG2 levels in all of the cattle inoculated with ISCOMATRIX/rMSPs (G1), compared to the control groups (G2 and G3) (p < 0.05). These data are in agreement with the results of a previous study, which found high IgG2 titers in cattle immunized with *A. marginale* outer membrane proteins (BROWN et al., 1998). The results obtained herein suggest that a mixed profile of immune response might occur in vivo after inoculation with rMSPs.

ISCOMATRIX/rMSPs induced the production of specific antibodies to each rMSP, as demonstrated by Western blot (Figure 3). Similar results were found when BALB/c mice were immunized with the same recombinant proteins (KAWASAKI et al., 2007b). In that study, it was suggested that a subunit vaccine containing rMSPs could be efficient in initial infection with *A. marginale*. Major surface protein 1a (MSP1a) and MSP1b occur as naturally complexed OMPs in the *A. marginale* outer membrane, and T-cell epitopes from MSP1a bound to MSP1b induced higher IgG titers against MSP1b (MacMILLAN et al., 2008). Additionally, it has been widely reported that clearance of *A. marginale* is dependent on high titers of IgG2-specific antibodies in cattle (BROWN et al., 1998; PALMER et al., 1999).

**Conclusion**

In conclusion, ISCOMATRIX/rMSPs induced an antigen-specific humoral immune response against MSPs of *A. marginale*, with the production of high levels of total IgG, IgG1 and IgG2 antibodies in immunized calves. The next step is to evaluate the protection afforded by rMSPs, including the use of new proteins previously detected as sub-dominant antigens, individually and collectively associated with adjuvant ISCOMATRIX against *A. marginale* strain challenges.

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